## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings of claims in the application:

## **Listing of Claims:**

- 1. (Original) A fission yeast strain comprising non-functional dgal and plh1 genes.
- 2. (Original) The yeast strain of claim 1 that is a Schizosaccharomyces pombe  $\Delta dgal \Delta plhl$  double deletion mutant.
- 3. (Presently Amended) The yeast strain of claim 1 or claim 2 comprising an exogenous gene that, when expressed in the *Schizosaccharonayces* yeast strain, results in TAG synthesis.
- 4. (Original) The yeast strain of claim 3 wherein the exogenous gene is a diacylglycerol acyl-transferase gene.
- 5. (Original) The yeast strain of claim 4 wherein the diacylglycerol acyltransferase gene is a human diacylglycerol acyltransferase gene.
- 6. (Original) A method for screening or identifying a compound that inhibits or prevents TAG synthesis, comprising: treating with a compound a culture of a fission yeast strain comprising non-functional *dgal* and *plhl* genes, wherein the yeast strain comprises an exogenous gene which is expressible in the yeast strain and which, when expressed in the yeast strain, results in TAG synthesis; and detecting any TAG synthesis in the culture.
- 7. (Original) The method of claim 6 wherein the yeast strain is a Schizosaccharomyces pompe  $\Delta dgal \Delta plhl$  double deletion mutant.
- 8. (Presently Amended) The method of claim 6 or claim 7 wherein the exogenous gene is a diacylglycerol acyl-transferase gene.

- 9. (Original) The method of claim 8 wherein the diacylglycerol acyltransferase gene is a human diacylglycerol acyltransferase gene.
- 10. (Presently Amended) The method of <u>claim 6</u> any one of <u>claims 6 to 9</u> wherein the compound is a small molecule, a protein, a peptide, an antibody, a hormone, a lipid or a nucleic acid.
- 11. (Presently Amended) The method of <u>claim 6</u> any one of claims 6 to 10 wherein the compound is useful for treatment of obesity, diabetes, coronary heart disease, heart failure or cardiomyopathy.
- 12. (Presently Amended) The method of <u>claim 6</u> any one of claims 6 to 11 wherein the detecting comprises adding labeled substrate of TAG synthesis to the culture.
- 13. (Original) The method of claim 12 wherein the substrate is labeled with a radioactive molecule, a chemiluminescent molecule, a fluorescent molecule, an enzyme that cleaves a reagent to produce a coloured molecule, a coloured molecule or a heavy metal complex.
- 14. (Presently Amended) The method of claim 12 or claim-13 wherein the labeled substrate is a fatty acid.
- 15. (Original) The method of claim 14 wherein the fatty acid is oleic acid or palmitic acid.
- 16. (Presently Amended) The method of <u>claim 12</u> any one of claims 12 to 15 wherein the detecting comprises extraction of cellular lipids and separation of the cellular lipids by thin layer chromatography.
- 17. (Presently Amended) The method of <u>claim 6</u> any one of claims 6 to 11 wherein the detecting comprises exposing the culture to conditions that are suitable for inducing

lipoapoptosis in a culture not expressing the exogenous gene and detecting lipoapoptosis in the exposed culture.

- 18. (Original) The method of claim 17 wherein the exposing comprises addition of fatty acid or diacylglycerol to the culture or to nutrient starvation.
- 19. (Original) The method of claim 18 wherein the fatty acid or diacylglycerol is added to a liquid culture during log phase.
- 20. (Presently Amended) The method of claim 17 or claim 18 wherein the fatty acid is oleic acid or palmitic acid and the diacylglycerol is diC8 diacylglycerol.
- 21. (Original) The method of claim 20 wherein the palmitic acid is added at a concentration of about 1 mM.
- 22. (Original) The method of claim 18 wherein the nutrient starvation comprises culturing the culture in water or low-glucose medium.
- 23. (Presently Amended) The method of <u>claim 17</u> any one of claims 17 to 22 wherein the detecting lipoapoptosis comprises measuring cell viability.
- 24. (Presently Amended) The method of <u>claim 17</u> any one of claims 17 to 23 wherein the detecting lipoapoptosis comprises detecting an apoptotic marker.
- 25. (Original) The method of claim 24 wherein the apoptotic marker is fragmented nuclear DNA, exposed phosphatidyl serine at the outer leaflet of the plasma membrane or production of reactive oxygen species.
- 26. (Presently Amended) The method of claim 24 [[or 25]] wherein the detecting lipoapoptosis comprises adding a detection molecule.
- 27. (Original) The method of claim 26 wherein the detection molecule is a radioactive molecule, a chemiluminescent molecule, a fluorescent molecule, an enzyme that

cleaves a reagent to produce a coloured molecule, a coloured molecule or a heavy metal complex.

- 28. (Original) A method of screening or identifying a compound that inhibits or prevents lipotoxicity, comprising: treating with a compound a culture of a fission yeast strain comprising non-functional *dgal* and *plhl* genes; exposing the treated culture to conditions that are suitable for inducing lipotoxicity in an untreated culture; and detecting lipotoxicity in the treated culture.
  - 29. (Original) The method of claim 28 wherein lipotoxicity is lipoapoptosis.
- 30. (Presently Amended) The method of claim 28 or claim 29 wherein the yeast strain is a *Sclaizosaccharoomyces pombe*  $\Delta dgal \Delta plhl$  double deletion mutant.
- 31. (Presently Amended) The method of <u>claim 28 any one of claims 28 to 30</u> wherein the compound is a small molecule, a protein, a peptide, an antibody, a hormone, a lipid or a nucleic acid.
- 32. (Presently Amended) The method of <u>claim 28</u> any one of claims 28 to 31 wherein the compound is useful for treatment of obesity, diabetes, coronary heart disease, heart failure or cardiomyopathy.
- 33. (Presently Amended) The method of <u>claim 28</u> any one of claims 28 to 32 wherein the exposing the treated culture comprises addition of fatty acid or diacylglycerol to the culture or nutrient starvation.
- 34. (Original) The method of claim 33 wherein the fatty acid or diacylglycerol is added to a liquid culture during log phase.
- 35. (Presently Amended) The method of claim 33 or claim 34 wherein the fatty acid is oleic acid or palmitic acid and the diacylglycerol is diC8 diacylglycerol.

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- 36. (Original) The method of claim 35 wherein the palmitic acid is added to a concentration of about 1 mM.
- 37. (Original) The method of claim 36 wherein the nutrient starvation comprises culturing the treated culture in water or low-glucose medium.
- 38. (Presently Amended) The method of <u>claim 29</u> any one of claims 29 to 37 wherein the detecting comprises detecting an apoptotic marker.
- 39. (Original) The method of claim 38 wherein the apoptotic marker is fragmented nuclear DNA, exposed phosphatidyl serine at the outer leaflet of the plasma membrane or production of reactive oxygen species.
- 40. (Presently Amended) The method of claim 38 [[or 39]] wherein the detecting lipoapoptosis comprises adding a detection molecule.
- 41. (Original) The method of claim 40 wherein the detection molecule is a radioactive molecule, a chemiluminescent molecule, a fluorescent molecule, an enzyme that cleaves a reagent to produce a coloured molecule, a coloured molecule or a heavy metal complex.
- 42. (Presently Amended) The method of <u>claim 29</u> any one of claims 29 to 37 wherein the detecting comprises measuring cell viability.
- 43. (Original) The method of claim 42 wherein measuring cell viability comprises performing a colony forming assay.
- 44. (Original) A method of making a fission yeast strain comprising non-functional dgal and plhl genes, comprising functionally interrupting the dgal and plhl genes in a fission yeast strain.
- 45. (Original) A method of screening or identifying a gene that complements non-functional dgal and plhl genes, comprising transforming a fission yeast strain comprising

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non- functional *dgal* and *plhl* genes with an exogenous gene; culturing the transformed yeast strain; and detecting any TAG synthesis in the culture.

- 46. (Original) A kit or commercial package comprising a fission yeast strain comprising non-functional *dgal* and *plhl* genes and instructions for screening or identifying a compound that inhibits or prevents TAG synthesis.
- 47. (Original) A kit or commercial package comprising a fission yeast strain comprising non-functional *dgal* and *plhl* genes and instructions for screening or identifying a compound that inhibits or prevents lipotoxicity.
- 48. (Original) A kit or commercial package comprising a fission yeast strain comprising non-functional *dgal* and *plhl* genes and instructions for screening or identifying a gene that complements the *dgal* and *plhl* genes.